

Notic of References Cited	Application/Control No.	Applicant(s)/Patent Under Reexamination ROULEAU ET AL.
	Examiner Joseph Woitach	Art Unit 1632

U.S. PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Name	Classification
	A	US-			
	B	US-			
	C	US-			
	D	US-			
	E	US-			
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	K	US-			
	L	US-			
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FOREIGN PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Country	Name	Classification
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NON-PATENT DOCUMENTS

*		Include as applicable: Author, Title Date, Publisher, Edition or Volume, Pertinent Pages)
	U	Grewal et.al.; Genetic mapping and haplotype analysis of oculopharyngeal muscular dystrophy, 1998, Neuro-Report 9: 961-965.
	V	Stajich et.al.; Confirmation of linkage of oculopharyngeal....existence of a second causal mutation, 1997, Neuromuscular Disorders 7: S75-S81.
	W	Akarsu et.al.; Genomic structure of HOXD13 gene: a nine polyalanine duplication causes synpolydactyly in two unrelated families, 1996, Human Molecular Genetics, Vol 5: 945-952.
	X	Brais et.al.; Oculopharyngeal Muscular Dystrophy, 1999, Seminars in Neurology, Vol. 19: 59-66.

*A copy of this reference is not being furnished with this Office action. (See MPEP § 707.05(a).)
Dates in MM-YYYY format are publication dates. Classifications may be US or foreign.

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	U	Brais et.al.; Short GCG expansions in the PABP2 gene cause oculopharyngeal muscular dystrophy, 1999, Nature Genetics, Vol.18: 164-167.
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